Roll Call No
Ayes
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Noes

HOUSE MOTION

MR. SPEAKER:

I move that House Bill 1857 be amended to read as follows:

1	Page 3, line 27, delete "." and insert "and any cost offsets or cost
2	shifts as a result of the cost containment measures."
3	Page 7, between lines 24 and 25, begin a new paragraph and insert:
4	"(e) The board may not require prior approval of a single source
5	drug based solely on the cost of the drug.
6	(f) The use of prior authorization must be based on the
7	recommendation of the board.".
8	Page 7, line 31, delete "shall" and insert "may".
9	Page 7, line 32, after "drug" delete "formulary and" and insert
10	"formulary. If the office establishes a formulary,".
11	Page 7, line 32, delete "board." and insert "board in compliance
12	with 42 U.S.C. 1396r.".
13	Page 9, between lines 15 and 16, begin a new paragraph and insert:
14	(I) A Medicaid managed care organization may not require
15	prior approval of a single source drug based solely on the cost of
16	the drug.
17	(m) The use of prior authorization must be based on the
18	recommendation of the board.
19	SECTION 14. IC 12-15-35-47, AS ADDED BY P.L.231-1999,
20	SECTION 9, IS AMENDED TO READ AS FOLLOWS [EFFECTIVE
21	JULY 1, 2001]: Sec. 47. (a) This section applies to the following
22	changes to a formulary used by a Medicaid managed care organization
23	for Medicaid recipients:
24	(1) Removing one (1) or more drugs from the formulary.

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1	(2) Placing new restrictions on one (1) or more drugs on the
2	formulary.
3	(b) Before a Medicaid managed care organization makes a change
4	described in subsection (a), the managed care organization shall submit
5	the proposed change to the office.
6	(c) The office shall forward the proposed change to the board for the
7	board's review and recommendation.
8	(d) The office shall provide at least thirty (30) days notification to
9	the public that the board will:
10	(1) review the proposed change; and
11	(2) consider evidence and credible information provided to the
12	board;
13	at the board's regular board meeting before making a recommendation
14	to the office regarding whether the proposed change should be
15	approved or disapproved.
16	(e) Based on the final recommendation of the board, the office may
17	approve or disapprove the proposed change. If a proposed change is not
18	disapproved within ninety (90) days after the date the managed care
19	organization submits the proposed change to the office, the managed
20	care organization may implement the change to the formulary.
21	(f) A Medicaid managed care organization:
22	(1) may add a drug to the managed care organization's formulary
23	without the approval of the office; and
24	(2) shall notify the office of any addition to the managed care
25	organization's formulary within thirty (30) days after making the
26	addition.
27	(g) A Medicaid managed care organization may not require
28	prior approval of a single source drug based solely on the cost of
29	the drug.
30	(h) The use of prior authorization must be based on the
31	recommendation of the board.".
32	Page 11, between lines 22 and 23, begin a new paragraph and insert:
33	"(e) Every six (6) months, the drug utilization review board
34	established by IC 12-15-35-19 shall:
35	(1) recommend to the office of Medicaid policy and planning
36	established by IC 12-15-1-1 those brand name drugs with
37	generic equivalents that are to be subject to prior approval;
38	and
39	(2) analyze:
40	(A) the cost savings achieved by the prior approval
41	program; and
42	(B) any concerns with:
43	(i) cost shifting; and
44	(ii) lowered health outcomes;
45	as a result of the prior approval program."
46	Renumber all SECTIONS consecutively.

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(Reference is to HB 1857 as printed February 16, 2001.)

Representative Crawford

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